

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Artcle 36 and Rule 70)

Applicant's or agent's file reference YL03002PCT	FOR FURTHER ACTION	SeeNotificationofTransmittalofInternationalPreliminary Examination Report (Form PCT/IPEA/416)		
International application No. PCT/KR2003/002034	International filing date(day/mo		Priority date (day/month/yell 11 OCTOBER 2002 (11.1	
International Patent Classification (IPC) IPC7 C07D 413/10	or national classification and IP	2		:
B & C BIOPHARM CO., LTD	. et al			
amended and are the basis for	according to Article 36.	ding this cover she of the description taining rectification	eet. n, claims and/or drawings w	hich have been
These annexes consist of a total of		·		
IV Lack of unity of inverse V X Reasoned statement citations and explana VI Certain documents c	of opinion with regard to novelty ention under Article 35(2) with regard tons supporting such statement	to novelty, inven		ability;
Date of submission of the demand	Date	of completion of	·	
11 MAY 2004 (11.	05.2004)	01 FEBRUAL	RY 2005 (01.02.2005)	
Name and mailing address of the IPEA/I Korean Intellectual Property 920 Dunsan-dong, Seo-gu, I Republic of Korea	Office Daejeon 302-701,	LEE, Jae Jeong	0.481.5404	ON O



INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International aplication No.

PCT/KR2003/002034

1.	Basis	s of the report	
1.	With	regard to the elements of the international application:*	
		the international application as originally filed	
	X	the description:	tot ono enca
		pages 12 pages	, as originally filed , filed with the demand
		pages 1 - 11, 13 , filed with the letter of 31/05/2	
	∇	the claims:	
	ب	pages, as amended (together with a	, as originally filed
		pages, as amended (together with a	, filed with the demand
		pages $14-15$, filed with the letter of $31/05/2$	2004
		the drawings:	
		pagespages	, as originally filed , filed with the demand
		pages, filed with the letter of	, med with the demand
		the sequence listing part of the description:	
		pagespages	, as originally filed
٠.		pages	_ , filed with the demand
2.	the i	n regard to the language, all the elements marked above were available or furnished to this Au international application was filed, unless otherwise indicated under this item.	
	Thes	se elements were available or furnished to this Authority in the following language	which is
		the language of a translation furnished for the purposes of international search (under Rule 2	23.1(b)).
		the language of publication of the international application(under Rule 48.3(b)).	
		the language of the translation furnished for the purposes of international preliminary examples of 55.3).	mination(under Rules 55.2 and/
3.		th regard to any nucleotide and/or amino acid sequence disclosed in the international application and the international application was carried out on the basis of the sequence listing:	plication, the international
		contained inthe international application in written form.	•
		filed together with the international application in computer readable form.	
		furnished subsequently to this Authority in written form.	
		furnished subsequently to this Authority in computer readable form	
		The statement that the subsequently furnished written sequence listing does not go international applicationas as filed has been furinshed.	beyond the disc losure in the
		The statement that the information recorded in computer readable form is identical to the	e written sequence listing has
•	Ш	been furnished.	•
		The second secon	
4.		The amendments have resulted in the cancellation of:	
	•	the description, pages	
		the claims, Nos.	
5.		the drawings, sheets	
<i>J</i> .		This report has been established as if (some of) the amendments had not been made, sing go beyond the disclosure as filed, as indicated in the Supplemental Box(Rule 70.2(c)).**	ce they have been considered to
	in thi	acement sheets which have been furnished to the receiving Office in response to an invitation is opinion as "originally filed." and are not annexed to this report since they do not contain 70.17).	
**	Any i	replacement sheet containing such amendments must be referred to under item I and annexed	l to this report.



International aplication No.

PCT/KR2003/002034

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement				
1. Statement				
Novelty (N)	Claims 1 - 3	YES		
	Claims	NO		
Inventive step (IS)	Claims 1 - 3	YES		
	Claims	NO		
Industrial applicability (IA)	Claims 1 - 3	YES		
	Claims	NO		

2. Citations and explanations (Rule 70.7)

Reference is made to the following documents:

D1: WO 0178648 A2 (Dong Wha Pharm. Ind. Co., Ltd.) 25 Oct. 2001

D2: EP 0187705 A2 (Norwich Eaton Pharmaceuticals, Inc.) 16 July 1986

D3: US 3878206 (Morton-Norwich Products, Inc.) 15 April 1975

D4: H.R. Snyder, et. al. 'Imidazo[4,5-f]quinolines III: Antibacterial 7-Methyl-9-(substituted Arylamino)imidazo[4,5-f]quinolines', In: Journal of Pharmaceutical Sciences, 1977, 66(8), pp.1204-6

The amended (31-05-2004) claims 1-3 of the present invention relates to 6-methylpyridine derivatives useful as an antiviral agent, and more particularly to novel 6-methylpyridine derivatives having an excellent inhibitory effect on replication of Hepatitis C virus (HCV), a method for preparing thereof, and an antiviral pharmaceutical composition comprising the compound as an active ingredient.

D1 discloses novel 6-methylnicotinamide derivatives, the process for preparing them, and the pharmaceutical compositions containing said compound as an active ingredient. The 6-methylnicotinamide derivatives of the present invention exhibit their inhibitory activity against the proliferation of HIV, HBV and HCV.

D2 concerns compounds of the class of imidazo[4,5-f]quinolines and methods for enhancing the immune response system of mammals which comprises systemically administering to mammals having a depressed immune function an effective but nontoxic amount of a composition comprising such a compound.

D3 and D4 describe a series of 9-(substituted amino)imidazo[4,5-f]quinolines as an antibacterial agent, particularly effective against Haemophilus vaginalis, a cause of bacterial vaginitis.

Although D1-D4 teach the process for preparing and using various types of 6-methylnicotinamide or imidazo[4,5-f]quinoline derivatives, D1-D4 do not disclose the features of the subject matter of claims 1 - 3, which meet the criteria set forth in PCT Article 33(2), (3) and (4). The 6-methylpyridine derivatives useful as an antiviral agent, a method for preparing thereof, and an antiviral pharmaceutical composition comprising the said compound are not anticipated by any of the references on record.

Thus, the invention described in the present application is considered to be novel, inventive and industrially applicable.





International application No. PCT/KR2003/002034

CLASSIFICATION OF SUBJECT MATTER IPC7 C07D 413/10 According to International Patent Classification (IPC) or to both national classification and IPC FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 07 C07D, A61K Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Korean Patents and Application for Inventions Since 1975 Electronic data base consulted during the intertnational search (name of data base and, where practicable, search terms used) CAS online(STN), Medline, Delphion DOCUMENTS CONSIDERED TO BE RELEVANT Category* Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. A WO 0178648 A2 (Dong Wha Pharm. Ind. Co., Ltd.) 25.Oct.2001 1-3 See whole document (Page 14, Compound 4) 1-3 EP 0187705 A2 (Norwich Eaton Pharmaceuticals, Inc.) 16.July 1986 See whole document US 3878206 (Morton-Norwich Products, Inc.) 15. April 1975 1-3 Α See whole document H.R. Snyder, Jr., C.F. Spencer and R. Freedman, Imidazo[4,5-f]quinolines III:Antibacterial 7-1-3 Α Methyl-9-(substituted Arylamino)imidazo[4,5-f]quinolines, Journal of Pharmaceutical Sciences, 1997, Vol.66(8), pp1204-6 See patent family annex. Further documents are listed in the continuation of Box C. Special categories of cited documents: "T" later document published after the international filing date or priority document defining the general state of the art which is not considered date and not in conflict with the application but cited to understand to be of particular relevance the principle or theory underlying the invention earlier application or patent but published on or after the international document of particular relevance; the claimed invention cannot be filing date considered novel or cannot be considered to involve an inventive document which may throw doubts on priority claim(s) or which is step when the document is taken alone cited to establish the publication date of citation or other document of particular relevance: the claimed invention cannot be special reason (as specified) considered to involve an inventive step when the document is document referring to an oral disclosure, use, exhibition or other combined with one or more other such documents, such combination being obvious to a person skilled in the art document published prior to the international filing date but later "&" document member of the same patent family than the priority date claimed Date of the actual completion of the international search Date of mailing of the international search report 09 JANUARY 2004 (09.01.2004) 10 JANUARY 2004 (10.01.2004) Authorized officer Name and mailing address of the ISA/KR Korean Intellectual Property Office 920 Dunsan-dong, Seo-gu, Daejeon 302-701, SIHN, YOUNG SIHN Republic of Korea

82-42-481-8162

Telephone No.

Facsimile No. 82-42-472-7140 .





INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No. PCT/KR2003/002034

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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